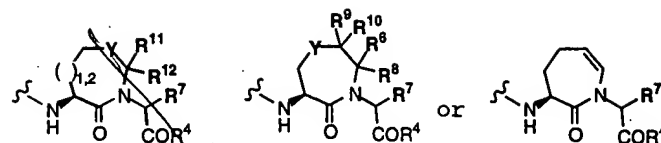


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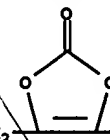
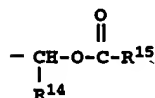
where Y is CH<sub>2</sub>,

R<sup>7</sup>, R<sup>8</sup> and R<sup>2</sup> are independently selected from hydrogen, alkyl, alkenyl, cycloalkyl-(CH<sub>2</sub>)<sub>m</sub>-, aryl-(CH<sub>2</sub>)<sub>m</sub>- and heteroaryl-(CH<sub>2</sub>)<sub>m</sub>-,

where m is 0 or an integer from 1 to 6;

R<sup>6</sup>, R<sup>10</sup>, R<sup>11</sup>, and R<sup>12</sup> are independently selected from hydrogen, alkyl, alkenyl, cycloalkyl-(CH<sub>2</sub>)<sub>p</sub>-, aryl-(CH<sub>2</sub>)<sub>p</sub>- and heteroaryl-(CH<sub>2</sub>)<sub>p</sub>-; and

R<sup>4</sup> is OH, alkyl, O-(CH<sub>2</sub>)<sub>p</sub>-heteroaryl,



-O-(CH<sub>2</sub>)<sub>p</sub>-aryl or -CH<sub>2</sub>-R<sup>16</sup> or NR<sub>1</sub>(R<sub>2</sub>) where R<sub>1</sub> and R<sub>2</sub> are independently H, alkyl, aryl, aryl-(CH<sub>2</sub>)<sub>p</sub> or heteroaryl;

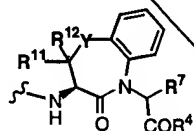
R<sup>14</sup> is hydrogen, alkyl, cycloalkyl or phenyl;

R<sup>15</sup> is hydrogen, alkyl, alkoxy or phenyl; and

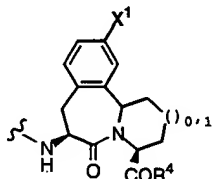
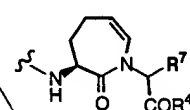
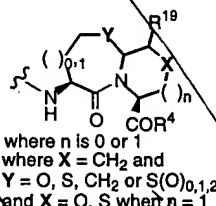
R<sup>16</sup> is alkyl or aryl-(CH<sub>2</sub>)<sub>m</sub>-.

6. (Amended) The compound as defined in Claim 1

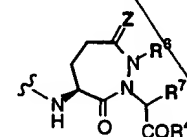
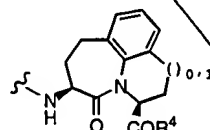
wherein A is



where Y = O, S, CH<sub>2</sub>, S(O)<sub>0,1,2</sub>



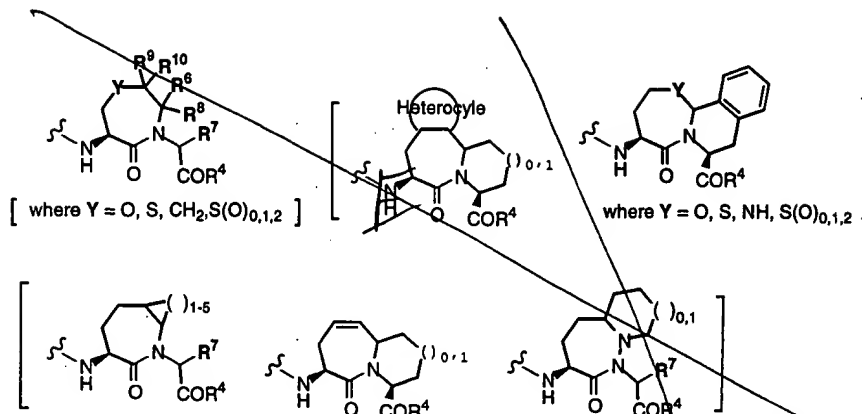
where X<sup>1</sup> = H, Ph,  
NHSO<sub>2</sub>R<sup>5</sup>  
(where R<sup>5</sup> ≠ H)



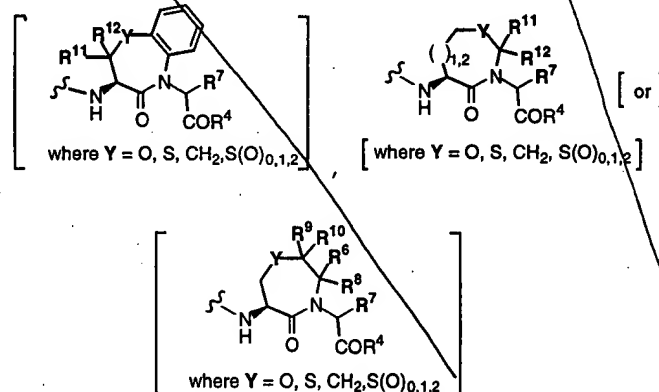
where Y = O, S, CH<sub>2</sub>, S(O)<sub>0,1,2</sub>  
where Z = O or H, H

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--7. (Amended) The compound as defined in Claim [6] 1  
wherein A is



--12. (Amended) A pharmaceutical composition comprising a  
[therapeutically effective amount of a] compound as defined in  
Claim 1 and a pharmaceutically acceptable carrier therefor.--

--13. (Amended) The pharmaceutical composition as  
defined in Claim 12 useful in the treatment of [cardiovascular  
diseases such as] hypertension and/or congestive heart failure.--

--14. (Amended) A method of treating [a cardiovascular  
disease such as] hypertension and/or congestive heart failure,  
which comprises administering to a mammalian species a  
therapeutically effective amount of a composition as defined in  
Claim 12.--

--15. (Amended) The compound as defined in Claim 1 which  
is

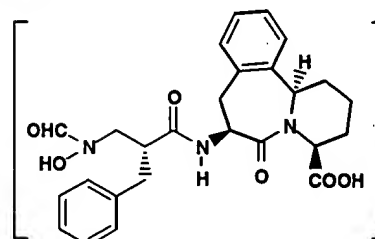
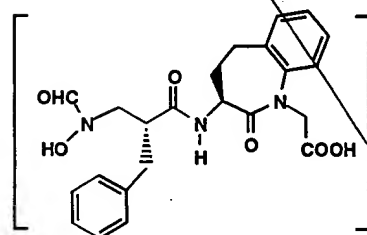
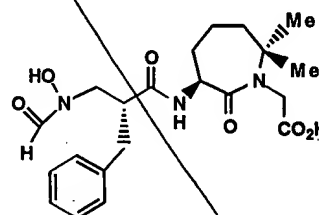
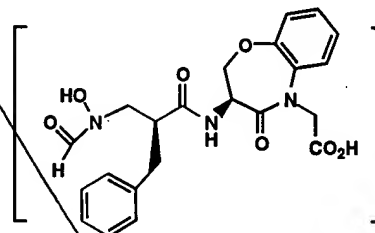
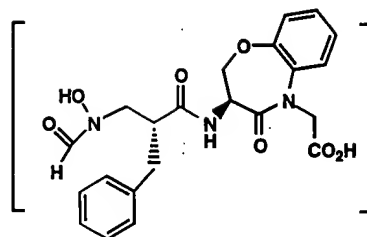
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or a pharmaceutically acceptable salt thereof.--